Complete Listing of the Claims:

Please cancel claims 1-38 and add new claims 39-87 as follows:

39. (New) A method for preparing a yeast mutant that produces a glycoprotein comprising a sugar chain represented by formula (IV), that comprises (i) introducing a GnT-I gene into a yeast mutant that produces a glycoprotein comprising a sugar chain represented by formula (III), and (ii) allowing a GnT-I protein to express therein, wherein formula III is represented by the following:

Man
$$\alpha$$
 1 6 Man α 1 6 Man α 1 6 Man α 1 7 3 Man β 1 – 4GlcNAc β 1 – 4GlcNAc (III)

wherein Man represents mannose and GlcNAc represents N-acetylglucosamine; and formula (IV) is represented by the following:

Man
$$\alpha$$
1 6 Man α 1 6 Man α 1 6 Man β 1 – 4GlcNAc β 1 – 4GlcNAc (IV) GlcNAc β 1 – 2Man α 1 / 3

wherein Man represents mannose and GlcNAc represents N-acetylglucosamine.

40. (New) A method for preparing a yeast mutant that expresses a GnT-I protein, an α-mannosidase II protein and a GnT-II protein, comprising introducing a GnT-I gene, an α-mannosidase II gene and a GnT-II gene into a yeast mutant that produces a glycoprotein comprising a sugar chain represented by the following formula

(III):
$$Man \alpha 1$$

wherein Man represents mannose and GlcNAc represents N-acetylglucosamine.

41. (New) The method according to claim 39, wherein said yeast mutant that produces a glycoprotein comprising a sugar chain represented by formula (IV) comprises an OCHI gene that is disrputed.

- 42. (New) The method according to claim 40, wherein said yeast mutant that expresses a GnT-I protein, an α-mannosidase II protein and a GnT-II protein comprises an OCH1 gene that is disrupted.
 - 43. (New) The method according to claim 39, wherein the yeast mutant is Saccharomyces cerevisiae.
- 44. (New) A method for preparing a yeast mutant that produces a glycoprotein comprising a sugar chain represented by formula (IV), which comprises (i) introducing an α -mannosidase I gene and a GnT-I gene into a yeast mutant that produces a glycoprotein comprising a sugar chain represented by formula (I), and (ii) allowing an α -mannosidase protein and a GnT-I protein to express therein, wherein formula (I) is represented by the following:

Man
$$\alpha$$
 1 – 2Man α 1 $\frac{6}{3}$ Man α 1 $\frac{6}{3}$ Man β 1 – 4GlcNAc β 1 – 4GlcNAc (I) Man α 1 – 2Man α 1 – 2Ma

wherein Man represents mannose and GlcNAc represents N-Acetylglucosamine; and formula (IV) is represented by the following:

Man
$$\alpha$$
 1 6 Man α 1 6 Man α 1 6 Man β 1 – 4GIcNAc β 1 – 4GIcNAc (IV)

wherein Man represents mannose and GlcNAc represents N-acetylglucosamine.

45. (New) A method for preparing a yeast mutant that expresses an α -mannosidase I protein, a GnT-I protein, an α -mannosidase II protein and a GnT-II protein, comprising introducing an α -mannosidase I gene, a GnT-I gene, an α -mannosidase II gene and a GnT-II gene into a yeast mutant that produces a glycoprotein comprising a sugar chain represented by the following formula (I):

Man
$$\alpha$$
 1 – 2Man α 1 $\frac{6}{3}$ Man α 1 $\frac{6}{3}$ Man α 1 – 4GlcNAc β 1 – 4GlcNAc (I)

Man α 1 – 2Man α 1 – 2Man α 1 $\frac{6}{3}$

wherein Man represents mannose and GlcNAc represents N-acetylglucosamine.

- 46. (New) The method according to claim 44, wherein said yeast mutant that produces a glycoprotein comprising a sugar chain represented by formula (IV) comprises an OCHI gene that is disrupted.
- 47. (New) The method according to claim 45, wherein said yeast mutant that expresses an α-mannosidase I protein, a GnT-I protein, an α-mannosidase II protein and a GnT-II protein comprises an OCHI gene that is disrupted.
- 48. (New) The method according claim 44, wherein the α -mannosidase I gene is derived from Aspergillus saitoi.
 - 49. (New) The method according to claim 44, wherein the yeast mutant is Saccharomyces cerevisiae.
 - 50. (New) The method according to claim 48, wherein the yeast mutant is Saccharomyces cerevisiae.
- 51. (New) A method for producing a yeast mutant that expresses an α-mannosidase I protein comprising introducing an α-mannosidase I gene into a yeast mutant that produces a glycoprotein comprising a sugar chain represented by the following formula (II).

Man
$$\alpha$$
 1 \sim 6

Man β 1 \sim 4GIcNAc β 1 \sim 4GIcNAc

Man α 1 \sim 2Man α

52. (New) A method for preparing a yeast mutant that expresses an α-mannosidase I protein, a GnT-I protein and a GnT-II protein, comprising introducing an α-mannosidase I gene, a GnT-I gene and a Gnt-II gene into a yeast mutant that produces a glycoprotein having a sugar chain represented by the following formula (II):

Man
$$\alpha$$
 1 \sim 6
Man β 1 – 4GIcNAc β 1 – 4GIcNAc
Man α 1 – 2Man α 1 – 2Man

53. (New) The method according to claim 51, wherein said yeast mutant that expresses an α-mannosidase I protein comprises an OCHI gene that is disrupted.

- 54. (New) The method according to claim 52, wherein said yeast mutant that expresses an α-mannosidase I protein, a GnT-I protein and a GnT-II protein comprises an OCHI gene that is disrupted.
- 55. (New) The method according to claim 51, wherein the α-mannosidase I gene is derviced from *Aspergillus saitoi*.
 - 56. (New) The method according to claim 51, wherein the yeast mutant is Saccharomyces cerevisiae.
 - 57. (New) The method according to claim 55, wherein the yeast mutant is Saccharomyces cerevisiae.
- 58. (New) A method for preparing a yeast mutant, which comprises disrupting with a regenerable auxotrophic marker in a yeast, a gene associated with a sugar chain synthesis of a glycoprotein produced by the yeast.
 - 59. (New) The method according to claim 58, wherein the auxotrophic marker is uracil.
- 60. (New) The method according to claim 58, wherein the gene associated with the sugar chain synthesis is an OCHI gene.
- 61. (New) A method for preparing a glycoprotein, which comprises (i) culturing the yeast mutant produced by the method according to claim 39 in a medium, (ii) producing and accumulating a glycoprotein in the obtained culture product, and (iii) collecting the glycoprotein from the culture product.
 - 62 (New) The glycoprotein produced by the method according to claim 39.
 - 63 (New) The method according to claim 40, wherein the yeast mutant is Saccharomyces cerevisiae.
 - 64. (New) The method according to claim 41, wherein the yeast mutant is Saccharomyces cerevisiae.
 - 65. (New) The method according to claim 42, wherein the yeast mutant is Saccharomyces cerevisiae.

- 66. (New) The method according claim 45, wherein the α-mannosidase I gene is derived from Aspergillus saitoi.
- 67. (New) The method according claim 46, wherein the α-mannosidase I gene is derived from Aspergillus saitoi.
- 68. (New) The method according claim 47, wherein the α-mannosidase I gene is derived from Aspergillus saitoi.
- 69. (New) The method according to any one of claims 45, wherein the yeast mutant is *Saccharomyces* cerevisiae.
- 70. (New) The method according to any one of claims 46, wherein the yeast mutant is Saccharomyces cerevisiae.
- 71. (New) The method according to any one of claims 47, wherein the yeast mutant is Saccharomyces cerevisiae.
 - 72. (New) The method according to claim 45, wherein the yeast mutant is Saccharomyces cerevisiae.
 - 73. (New) The method according to claim 46, wherein the yeast mutant is Saccharomyces cerevisiae.
 - 74. (New) The method according to claim 47, wherein the yeast mutant is Saccharomyces cerevisiae.
- 75. (New) The method according to claim 52, wherein the α-mannosidase I gene is derviced from *Aspergillus saitoi*.
- 76. (New) The method according to claim 53, wherein the α-mannosidase I gene is derviced from *Aspergillus saitoi*.
- 77. (New) The method according to claim 54, wherein the α-mannosidase I gene is derviced from Aspergillus saitoi.

- 78. (New) The method according to claim 52, wherein the yeast mutant is Saccharomyces cerevisiae.
- 79. (New) The method according to claim 53, wherein the yeast mutant is Saccharomyces cerevisiae.
- 80. (New) The method according to claim 54, wherein the yeast mutant is Saccharomyces cerevisiae.
- 81. (New) The method according to claim 75, wherein the yeast mutant is Saccharomyces cerevisiae.
- 82. (New) The method according to claim 76, wherein the yeast mutant is Saccharomyces cerevisiae.
- 83. (New) The method according to claim 77, wherein the yeast mutant is Saccharomyces cerevisiae.
- 84. (New) The method according to claim 69, wherein the gene associated with the sugar chain synthesis is an OCHI gene.
- 85. (New) The method according to claim 707, wherein the gene associated with the sugar chain synthesis is an OCHI gene.
- 86. (New) The method according to claim 71, wherein the gene associated with the sugar chain synthesis is an OCHI gene.
- 87. (New) The method according to claim 50, wherein the gene associated with the sugar chain synthesis is an OCHI gene.